Applicant:

Title:

Peripheral-Type Benzodiazepine Receptor: A Tool for Detection, Diagnosis, Prognosis, and Cancer

1941.016US2

Treatment of Cancer

Docket No.:

Filed:

September 25, 2000

Examiner:

Unknown

Commissioner for Patents

P.O. Box 1450

Alexandria, VA 22313-1450

Due Date: April 18, 2008 Group Art Unit: 1642

We are transmitting herewith the following attached items (as indicated with an "X"):

Return postcard.

Preliminary Amendment (8 pgs.).

Sequence Listing (2 pages) with computer readable form (1 diskette).

A Copy of the Notice to Comply with Requirement for Patent Applications containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures as mailed on March 18, 2008 (1 pg.).

Please consider this a PETITION FOR EXTENSION OF TIME for sufficient number of months to enter these papers and please charge any additional fees or credit overpayment to Deposit Account No. 19-0743.

SCHWEGMAN, LUNDBERG & WOESSNER, P.A.

Customer Number 21186

CERTIFICATE UNDER 37 CFR 1.8: The undersigned hereby certifies that this correspondence is being deposited with the United States Postal Service with sufficient postage as first class mail, in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450, on this 17th day of April, 2008.

Name

SCHWEGMAN, LUNDBERG & WOESSNER, P.A.

(GENERAL)



Application No.: 09/646932

4.

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. § 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

1. This application clearly fails to comply with the requirements of 37 C.F.R. 1821-1825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 11 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).
induce published at 65 FR/29020 (June 1, 1998) and 1211 OG 82 (June 23, 1998).
2 This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence" Listing as required by 37 C.F.R. 1.821(c).
3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by
37 C.F.R. 1821(e)
4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822
and/or 1 823, as indicated on the attached copy of the marked -up "Raw Sequence Listing"
5. The computer readable form that has been filed with this application has been found to be damaged
and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d)
6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the
"Sequence Listing" as required by 37 C.F.R. 1.821(e).
7. Other:
Applicant Must Provide:
An initial or substitute computer readable form (CRF) copy of the "Sequence Listing"
An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its en
into the specification.
A statement that the content of the paper and computer readable copies are the same and whereave applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).
For questions regarding compliance to these requirements, please contact
For Rules Interpretation, call (703) 308-4216
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